

detection systems; and training personnel in Privacy Act and information security requirements. After the records have been scheduled with NARA, records that are eligible for destruction will be disposed of in accordance with the applicable schedule, using secure destruction methods prescribed by NIST SP 800–88.

RECORD ACCESS PROCEDURES:

An individual seeking access to records about that individual in this system of records must submit a written access request to the applicable System Manager identified in the “System Manager” section of this System of Records Notice (SORN). The request must contain the requester’s full name, address, and signature. The request should also contain sufficient identifying particulars (such as, the provider’s National Provider Identifier, TIN, or patient medical record number, or the patient’s patient identifier or SSN) to enable HHS to locate the requested records. So that HHS may verify the requester’s identity, the requester’s signature must be notarized or the request must include the requester’s written certification that the requester is the individual who the requester claims to be and that the requester understands that the knowing and willful request for or acquisition of a record pertaining to an individual under false pretenses is a criminal offense subject to a fine of up to \$5,000.

If an access request by a patient seeks disclosure of any information about the patient’s provider which is or could be proprietary information of that provider, the request must be accompanied by a disclosure authorization form signed by the provider.

CONTESTING RECORD PROCEDURES:

An individual seeking to amend a record about that individual in this system of records must submit an amendment request to the applicable System Manager identified in the “System Manager” section of this SORN, containing the same information required for an access request. The request must include verification of the requester’s identity in the same manner required for an access request; must reasonably identify the record and specify the information contested, the corrective action sought, and the reasons for requesting the correction; and should include supporting information to show how the record is inaccurate, incomplete, untimely, or irrelevant.

NOTIFICATION PROCEDURES:

An individual who wishes to know if this system of records contains records about that individual should submit a notification request to the applicable System Manager identified in the “System Manager” section of this SORN. The request must contain the same information required for an access request and must include verification of the requester’s identity in the same manner required for an access request.

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

None.

[FR Doc. 2021–25760 Filed 11–24–21; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; The Stem Cell Therapeutic Outcomes Database, OMB No. 0915–0310—Extension

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than January 25, 2022.

ADDRESSES: Submit your comments to paperwork@hrsa.gov or by mail to the HRSA Information Collection Clearance Officer, Room 14N136B, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email paperwork@hrsa.gov or call Samantha Miller, the acting HRSA Information Collection Clearance Officer at (301) 443–9094.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the

information collection request title for reference.

Information Collection Request Title: The Stem Cell Therapeutic Outcomes Database OMB No. 0915–0310—Extension

Abstract: Given the rapid evolution of COVID–19 and its impact on those with compromised immune systems, it is imperative for the transplant community to continue collecting COVID–19 related data. Having access to COVID–19 vaccination status on blood stem cell recipients and understanding immune responses will assist with making informed decisions regarding direct clinical care. This will also inform critical policy decisions.

The Stem Cell Therapeutic and Research Act of 2005, Public Law (P.L.) 109–129, as amended, provides for the collection and maintenance of human blood stem cells for the treatment of patients and research. It also maintains a scientific database of information relating to patients who have been recipients of a stem cell therapeutics product (e.g., bone marrow, cord blood, or other such product) from a donor.

Given the rapid evolution of the COVID–19 public health emergency and its impact on immunocompromised patients, availability of new vaccines, and continual changes in vaccination recommendations, HRSA wants to leverage the required data collection platform of the Stem Cell Therapeutic Outcomes Database to obtain vaccine information for all U.S. allogeneic hematopoietic stem cell transplant recipients.

Need and Proposed Use of the Information: To collect COVID–19 vaccine data, HRSA is requesting an extension of OMB’s approval of both the Pre-Transplant Essential Data Form 2400 and Post-Transplant Essential Data (Post-TED) Form 2450. Collecting these data will help clinicians and policymakers to understand the landscape of vaccination among immunocompromised patients before and after a blood stem cell transplant.

HRSA will use this information to analyze outcomes based on vaccine manufacturer/type, doses received (including potential boosters), timing, and inform future vaccination strategies. Information currently collected regarding COVID–19 infections has already been used in research studies.

HRSA will use data collected prior to a patient receiving a blood stem cell transplant to characterize frequencies of vaccination and level of protection afforded during and after transplant based on incidence of COVID infection. Post-transplant, this information can be used to assess vaccination rates and

timing in blood stem cell recipients, characterize emerging vaccination strategies (which may include boosters), describe possible short and long-term side effects of vaccines, and analyze the incidence of COVID-19 infection based on different vaccination approaches. This information may guide future vaccination strategies or COVID treatments. Vaccination status of recipients may also be useful for risk adjustment in the annual transplant center specific analysis. For example, Centers for Disease Control and Prevention advisors could potentially use COVID-19 vaccination data on blood stem cell transplant recipients to make informed decisions regarding whether to issue any recommendations for this medically vulnerable population. The data collected under this extension request could help answer these and other questions.

The additional COVID-19 vaccine questions capture basic information on vaccination status, vaccine manufacturer/type, dose(s) given, and

date(s) received. Patients who need a blood stem cell transplant are typically aware of their COVID-19 risk and vaccination status, and the information is also found on the vaccine cards carried by most recipients. Questions about vaccination status will likely become universal during the intake process at transplant centers for the next 12 months or more. For these reasons, HRSA believes the data will be readily available to data professionals working at transplant centers via the medical record. To reduce burden, an “unknown” option has been included for scenarios where the data cannot be located, and a “date estimated” checkbox has been included when the exact date of vaccination is not known. Although these questions are anticipated to be asked over the next 12 months and then removed, it is possible that other COVID-19 related questions may be requested for inclusion on these forms in the future given the rapid evolution of COVID-19 and its impact on immunocompromised patients,

availability of new vaccines, and continual changes in vaccination recommendations.

Likely Respondents: Transplant Centers.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents ¹	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Baseline Pre-Transplant Essential Data (TED)	200	48	9,600	² 0.70	6,720
Disease Classification	200	48	9,600	³ 0.43	4,160
Product Form (includes Infusion, HLA, and Infectious Disease Marker inserts)	200	45	9,000	1.00	9,000
100-day Post-TED	200	48	9,600	0.88	8,448
6 month Post-TED	200	43	8,600	0.85	7,310
1 year Post-TED	200	40	8,000	0.65	5,200
2 year Post-TED	200	34	6,800	0.65	4,420
3+ years Post-TED	200	172	34,400	⁴ 0.52	17,773
Total	200		95,600		63,031

¹ The total of 200 is the number of centers completing the form; the same group will complete all of the forms.

² The decimal is rounded down, and the actual number is .683333333.

³ The decimal is rounded down, and the actual number is .433333333.

⁴ The decimal is rounded up, and the actual number is .516667.

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Maria G. Button,

Director, Executive Secretariat.

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BILLING CODE 4165-15-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Federal Financial Participation in State Assistance Expenditures; Federal Matching Shares for Medicaid, the Children's Health Insurance Program, and Aid to Needy Aged, Blind, or Disabled Persons for October 1, 2022 Through September 30, 2023

AGENCY: Office of the Secretary, DHHS.
ACTION: Notice.

The Federal Medical Assistance Percentages (FMAP), Enhanced Federal Medical Assistance Percentages (eFMAP), and disaster-recovery FMAP adjustments for Fiscal Year 2023 have been calculated pursuant to the Social

Security Act (the Act). These percentages will be effective from October 1, 2022 through September 30, 2023. This notice announces the calculated FMAP rates, in accordance with sections 1101(a)(8) and 1905(b) of the Act, that the U.S. Department of Health and Human Services (HHS) will use in determining the amount of federal matching for state medical assistance (Medicaid), Temporary Assistance for Needy Families (TANF) Contingency Funds, Child Support Enforcement collections, Child Care Mandatory and Matching Funds of the Child Care and Development Fund, Title IV-E Foster Care Maintenance payments, Adoption Assistance payments and Kinship Guardianship